Dermatological manifestations of COVID-19 and its therapies: a review

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INTRODUCTION

SARS-CoV2 (severe acute respiratory syndrome coronavirus-2), a novel coronavirus was first reported in December of 2019 from Wuhan, China, as an etiological agent causing a new infectious respiratory disease (coronavirus disease 2019, COVID-19).1

SARS-CoV-2 spread rapidly worldwide through human-to-human transmission following which COVID-19 was declared as a pandemic emergency by world health organization (WHO) on March, 2020.

By October 14th 2020, 38,363,705 patients were confirmed as COVID-19 cases and total toll of death reaching up to 1,090,811 patients.

The main clinical manifestations of COVID-19 being fever, cough, fatigue, dyspnoea and muscle aches.2,3 The diarrhoea, olfactory and gustatory impairments are some recent symptoms to join the list occurring in 5-15% of patients.4,5 The skin being the largest organ of the body was not very far from becoming prey to COVID19 virus.

The main dermatological manifestations were classified under 5 classes as of-
Erythematous rash: Erythematous rash (47%) (morbilliform, maculopapular, papulosquamous). Generally, appear within 6 days of appearance of first symptom of COVID-19 and involve legs, thighs, forearms, arms, shoulders, back, chest and abdomen.6

Urticarial eruptions: Urticarial eruptions (19%) pruritic wheals and plaques, appear on 2-5 days of disease and involve entire body while in case of urticarial vasculitis involvement of trunk and limbs is seen.7

Varicella-like vesicular eruptions: Nine percent chicken pox like vesicles on an erythematous background predominantly appear on trunk and limbs with mean duration of appearance being 5 days of disease onset.8

Acral lesions (COVID toes): Nineteen percent acral ischemia leads to formation of pseudochilblains like lesions over distal extremities (hands and feet) with mean duration appearance being 8 days.9

Livedoid eruptions: Livedoid eruptions (6%) a retiform pigmentation predominantly over lower extremities seldom breaking into ulcers which heal onto give porcelain white scarring presents on 1-3 days of disease onset.10

Recently two more classes acrocyanosis bluish discoloration of fingers and toes which appears as early as third day of disease onset and Retiform purpura appearing on 5-15 days of disease onset predominantly involving buttocks, back and abdomen.11,12

Were added to the pre-existing list of symptoms along with many other isolated cases of different cutaneous features.

Based on the evidence published to date, the cutaneous manifestations of novel coronavirus are similar to those caused by other common viral infections, all owing to their mostly similar pathogenesis.

The rashes (morbilliform, papulosquamous) characteristic of the acute phase of the inflammatory response generated by viral infection, could appear early in infection. Whereas Acral lesions and/or chilblain-like lesions (COVID toes) could be a late manifestation of inflammatory processes or microthrombotic events in the immune phase of disease.13 Some studies tried to establish a temporal relationship between skin lesions and the severity of systemic symptoms. For e.g.- pseudo-chilblain was associated with less severe pulmonary disease in contrast to livedoid presentations which were associated with worse pulmonary involvement.14

CLASSIFICATION OF SKIN LESIONS DUE TO SARS COV2

After reviewing many articles and case reports on dermatological manifestations of COVID-19 worldwide, we got the understanding that some clinical features were more commonly found, while some were individual presentations. Hence, we decided to categorize all the features into established presentations and isolated presentations in their respective age groups (Table 1).15-25

Table 1: Classification of dermatological manifestations in COVID-19 patients in their respective age groups.

<table>
<thead>
<tr>
<th>Adult Established</th>
<th>Adult Isolated</th>
<th>Paediatric Established</th>
<th>Paediatric Isolated</th>
</tr>
</thead>
<tbody>
<tr>
<td>Erythematous rash- Morbilliform rash, Maculopapular rash, papulosquamous rash</td>
<td>Symmetrical drug-related intertriginous and flexural exanthema (SDRIFE)-like erythematous rash</td>
<td>Hand-foot-mouth disease</td>
<td>Acute haemorrhagic edema of infancy</td>
</tr>
<tr>
<td>Urticaria and Urticarial vasculitis</td>
<td>Idiopathic plantar hidradenitis</td>
<td>Chickenpox like vesicles</td>
<td>Ulcerated facial skin</td>
</tr>
<tr>
<td>Vesiculobullous rashes-Variella-like papulovesicular exanthema</td>
<td>Neutrophilic dermatosis</td>
<td>Varicella-like exanthem</td>
<td></td>
</tr>
<tr>
<td>Acral ischemia-Lupus erythematosus (LE)-like chilblains of feet (COVID toes)</td>
<td>Acute acral cyanosis</td>
<td>P. Rosea like lesions</td>
<td></td>
</tr>
<tr>
<td>COVID-19-related vasculitis-Livedo reticularis</td>
<td>COVID red half-moon nail sign</td>
<td>Maculopapular rash</td>
<td></td>
</tr>
<tr>
<td>Erythema multiforme like lesions</td>
<td>Dengue like Petechial rash</td>
<td>Kawasaki disease</td>
<td></td>
</tr>
<tr>
<td>Retiform purpura</td>
<td>Androgenetic alopecia (AGA)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Acute generalized exanthematous pustulosis (AGEP)-like rash</td>
<td>Pigment disorders-Diffuse melanoderma of acute onset (peri orbital dyschromia)</td>
<td>Dry gangrene</td>
<td>Eosinophilic panniculitis</td>
</tr>
</tbody>
</table>
Histopathological findings in COVID-19 skin

In maculopapular eruptions: Epidermis shows dyskeratotic cells, ballooning multinucleated cells and sparse necrotic keratinocytes with lymphocytic satellitosis and nests of Langerhans cells. In Dermis-diffuse telangiectatic blood vessels, Perivascular spongiotic dermatitis and a dense perivascular lymphocytic infiltration eosinophilic rich around swollen blood vessels with extravasated erythrocytes are seen. Edematous dermis with many eosinophils, cuffs of lymphocytes around blood vessels in lymphocytic vasculitis and intravascular microthrombi in small dermal vessels is present. 26

In cases of urticaria like rash: Superficial perivascular inflammation with eosinophils and lichenoid pattern, and sometimes Slight vacuolar-type interface dermatitis with occasional necrotic keratinocytes may be seen.27

In cases of vasculitis: An inflammatory infiltrate in wall of dermal or subcutaneous vessels (which can be neutrophilic, lymphocytic, or granulomatous), red blood cell extravasation, variable fibrinoid necrosis of vessel walls, and leucocytoclasia is seen. Further on DIF staining, deposits of immunoglobulin, complement, or fibrin in the vessel wall can be seen.28

In case of vesicobullous eruptions: Vacuolar degeneration of the basal layer with multinucleate, hyperchromatic keratinocytes and dyskeratotic cells with mixed inflammatory infiltrate are present.29

In cases of livedo vasculopathy: Thrombogenic vasculopathy accompanied by extensive deposition of C5b-9 and C4d within the microvasculature are seen.30

In cases of acral perniosis: A diffuse dense lymphoid infiltrate of the superficial and deep dermis, and signs of endothelial activation (swelling and infiltration) with mild superficial perivascular dermatitis are present.31

Pathogenetic mechanism of dermatological manifestations in COVID-19

The pathological mechanisms of skin lesions in COVID-19 patients remain poorly understood.

Cutaneous manifestations in COVID-19 may be classified into two major groups regarding their pathomechanisms- (A) Clinical features similar to viral exanthems, due to direct effect of COVID19 virus. (B) Cutaneous eruptions secondary to systemic consequences caused by COVID-19 (especially vasculitis and thrombotic vasculopathy).32

SARS-CoV-2 is a single-stranded RNA virus composed of 16 non-structural proteins (NSP 1-16) with specific roles in the replication of coronaviruses.33 Where, NSP3 has the ability to block the host’s innate immune response and promote cytokine expression, while NSP5 can inhibit interferon (IFN) signalling, and NSP16 avoids MAD5 (melanoma differentiation-associated gene recognition, while depressing innate immunity.34

Different morphological lesions and pathogenesis implicated in their appearance- Vesicular lesions have been commonly described with COVID-19. Pathogenesis has been poorly understood; however, the presence of multinucleated ballooning cells suggests a direct cytopathic effect, lending weight to the hypothesis that lesions are due to COVID-19 virus related cytopathy.35

COVID-19 progression is related to an extreme rise in inflammatory cytokines including interleukin (IL)2, IL7, IL10, GCSF, IP10, MCP1, MIP1 A, and TNFα. The increase in the pro-inflammatory cytokines, in particular, IL6 is associated with severe pneumonia and it can have deleterious effects on the adaptive immune system.36 In these subsets of patients, overactive immune responses may induce immunopathological conditions, named as cytokine storm and in some individuals leads to macrophage activation syndrome (MAS)-like, often causing a fatal outcome.37 Cytokines could reach the skin and stimulate dermal dendritic cells, macrophages, mast cells and lymphocytes, in addition to polymorphonuclear cells and promote eruptions such as erythema, urticarial lesions, vesicles and others.

Hamming et al identified the metallopeptidase named angiotensin-converting enzyme 2 (ACE2) as the functional receptor for SARS-CoV responsible for an epidemic outbreak during 2003-2004.38 Using IHC methods, it was found that the surface expression of ACE2 protein was present on lung alveolar epithelial cells (pneumocytes), macrophages, enterocytes of the small intestine, arterial and venous endothelial cells, arterial smooth muscle cells including the skin in the basal layer of the epidermis, endothelial cells of dermal blood vessels and eccrine adnexal tissue.39 The ACE2 receptor is also widely expressed on endothelial cells in multiple organs, suggesting that endothelitis could occur in several sites as a direct consequence of viral involvement and host inflammatory response.

Varga et al revealed viral inclusion structures in endothelial cells across vascular beds of different organs in some patients with COVID-19.40 This showed direct viral infection of endothelial cell and diffuse endothelial inflammation. COVID-19-endothelitis could explain the systemic impaired microcirculatory function in different vascular beds and their clinical sequelae in patients with COVID-19. SARS-CoV2 binds to ACE2 by spike protein (S) and this allows entry of virus inside the cell.41 In order for virus to complete entry into the cell following this initial process, the spike protein has to be primed by a protease (TMPRSS2) to complete this process.42 The expression of this protease (TMPRSS2) is androgen dependent, and this maybe the reason why males are more commonly and severely affected by COVID-19.43
Complement activation (C5b-9 and C4d) by SARS-CoV-2 spike glycoproteins has been seen in retiform purpura. In pseudo chilblain and purpuric lesions, an obliterative microangiopathy consisting of endothelial and intensive myointimal growth with complement activation has been observed. This mechanism, together with increased vascular permeability, could contribute to obliterative vascular lumen and haemorrhage in COVID patients.44

A proper understanding of these pathomechanisms will help in formulating newer therapies for treating dermatological manifestations of COVID-19.

There has always been a diagnostic dilemma regarding the origin of cutaneous features in the course of COVID-19 infection, whether the dermatological signs are a result of COVID-19 infection or just an ADR to the drugs implicated in the treatment of patient. Keeping Proper timeline of the appearance of lesions and monitoring and correlating the severity of COVID symptoms with dermatological manifestation may help to rule out the underlying cause in some crossover cases.

Here we are discussing some common drugs advocated in treatment of COVID worldwide and summarising some common drug reactions reported by them (Table 2).45

Table 2: Drugs advocated in treatment of COVID-19 and cutaneous manifestation of ADR because of them.

<table>
<thead>
<tr>
<th>Drugs</th>
<th>Adverse effects</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Antimalarials-Hydroxychloroquine and chloroquine</strong></td>
<td>Acute generalized exanthematous pustulosis, Urticaria, pruritus, rashes Flares of psoriasis and exfoliating lesions, Stevens-Johnson-like syndrome Mucocutaneous dyspigmentation</td>
</tr>
<tr>
<td><strong>Azithromycin</strong></td>
<td>Generalized red or purple skin rashes Anaphylaxis DRESS syndrome Cutaneous leukocytoclastic vasculitis Fixed drug eruptions.</td>
</tr>
<tr>
<td><strong>Colchicine</strong></td>
<td>Blanchable violaceous morbilliform rash Lichenoid drug eruption Alopecia Toxic epidermal necrolysis-like reaction Erythema-nodosum-like lesions.</td>
</tr>
<tr>
<td><strong>Remdesivir</strong></td>
<td>Skin rashes</td>
</tr>
<tr>
<td><strong>Ivermectin</strong></td>
<td>Swelling of ankles and hands Allergic skin reactions such as Urticaria and Rash</td>
</tr>
<tr>
<td><strong>Oseltamivir</strong></td>
<td>Stevens-Johnson syndrome/toxic epidermal necrolysis, Angioedema Allergic or an idiosyncratic cutaneous drug reaction</td>
</tr>
<tr>
<td><strong>Ribavirin</strong></td>
<td>Acneiform eruptions, Alopecia Localized scleroderma Maculopapular lesions Skin dryness, Pruritus and rash</td>
</tr>
<tr>
<td><strong>Interferons</strong></td>
<td>Injection site reactions, Psoriasis Alopecia Sarcoïdosis Lupus Cutaneous vasculitis lesions Lichenoid drug reactions</td>
</tr>
<tr>
<td><strong>Protease inhibitors (ritonavir/lopinavir)</strong></td>
<td>Maculopapular drug eruptions Exfoliative erythroderma Stevens-Johnson syndrome or TEN Injection site reactions</td>
</tr>
<tr>
<td><strong>Corticosteroids</strong></td>
<td>Acneiform eruptions Folliculitis Skin atrophy Telangiectasia Hirsutism Stria</td>
</tr>
</tbody>
</table>

Continued.
<table>
<thead>
<tr>
<th>Drugs</th>
<th>Adverse effects</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Tocilizumab</strong></td>
<td>Skin hypersensitivity reactions, Psoriasiform dermatitis</td>
</tr>
<tr>
<td><strong>Anakinra</strong></td>
<td>Skin rashes  Injection-site reactions  Cellulitis</td>
</tr>
<tr>
<td><strong>JAK inhibitors (Baricitinib)</strong></td>
<td>Palmoplantar pustulosis-like eruption  Herpes zoster and simplex activations  Melanoma and nonmelanoma skin cancers  Urticaria, Rash, Angioedema</td>
</tr>
<tr>
<td><strong>Anti-TNF biologies (Adalimumab)</strong></td>
<td>Psoriasiform-like lesions  Lupus-like syndromes  Cutaneous vasculitis  Granulomatous reactions</td>
</tr>
<tr>
<td><strong>IVIG treatments</strong></td>
<td>Anaphylactic reaction  Facial vasculitis rash  Maculopapular rashes  Erythema multiforme  Alopecia</td>
</tr>
<tr>
<td><strong>Zinc supplementation</strong></td>
<td>Generalised dilution in skin color  Menkes kinky hair disease  Trichoptilosis</td>
</tr>
<tr>
<td><strong>Vitamin C</strong></td>
<td>Nil</td>
</tr>
</tbody>
</table>

**ANTIMALARIALS**

Hydroxychloroquine and chloroquine have antiviral activity against COVID-19 in vitro and small-uncontrolled clinical studies and has been the mainstay for treatment as well as prophylaxis for health care workers in the beginning of pandemic.

Cutaneous adverse events of antimalarials include cutaneous eruptions such as acute generalized exanthematous pustulosis, urticaria, pruritus, dry skin, rashes, flares of psoriasis and exfoliating lesions, Stevens-Johnson-like syndrome, mucocutaneous dyspigmentation, alopecia and bleaching of hair.45

**Azithromycin**

In the recent treatment guidelines issued by AIIMS New Delhi, azithromycin is indicated in all mildly symptomatic patients practicing home quarantine. Azithromycin has immunomodulatory and anti-inflammatory properties hence Combining hydroxychloroquine with the antibiotic azithromycin has been associated with positive patient outcomes according to a low-powered French study Skin adverse events of azithromycin are generalized red or purple skin rashes, blistering, skin peeling, toxic pustuloderma, anaphylaxis, DRESS syndrome, cutaneous leukocytoclastic vasculitis, and fixed drug eruptions.45

**Colchicine**

Colchicine is being used for its anti-neutrophilic immunomodulatory effect; it decreases the chances of severe pulmonary involvement, organ failure, and death which are due to cytokine storm and inflammatory process. Skin side-effects of colchicine include diffuse, blanchable, violaceous, morbilliform rash, lichenoid drug eruption, alopecia, toxic epidermal necrolysis-like reaction, erythema-bullous, and erythema-nodosum-like lesions.45

**Remdesivir**

It is being advocated in moderately symptomatic hospitalised patients of COVID-19 and with CT score >12 according to their recent CT scan chest. Remdesivir shuts down viral replication by inhibiting a key viral enzyme, the RNA polymerase. The drug, which is given intravenously, and the most common adverse events is skin rashes.45

**Oseltamivir**

This has also been approved for the treatment of pandemic influenza A and B treatments. Oseltamivir inhibits the viral neuraminidase and, blocks the release of viral particles from host cells. Oseltamivir can cause Stevens-Johnson syndrome/toxic epidermal necrolysis, angioedema, allergic, or idiosyncratic cutaneous drug reactions.45

**Favipiravir**

Favipiravir is a nucleoside analogue that is well-known as a broad-spectrum antiviral drug with a lower side-effects profile. Its oral preparation recently being marketed under the name of FABIFLU is being widely used in mildly
symptomatic cases of COVID-19. No significant data was collected regarding the cutaneous side effect of favipravir.\textsuperscript{45}

**Umifenovir**

inhibits membrane fusion of the viral envelope and spike S protein/ACE2 interaction. Basically, inhibiting the fusion and entry of virus inside the cells. There are no reports on the skin reaction of umifenovir.\textsuperscript{45}

**Ribavirin**

It is a broad antiviral against respiratory viruses such as influenza A and B viruses and parainfluenza 1 virus. Side effects of ribavirin include acneiform eruptions, alopecia, localized scleroderma, maculopapular, and eczematous lesions, skin dryness, pruritus, and rash.\textsuperscript{45}

**Interferon alpha-2a**

interact with the toll-like receptors and can inhibit the viral replication. Cutaneous side-effects of interferons are injection site reactions, psoriasis, eczematous drug reactions, alopecia, sarcoidosis, lupus, cutaneous vasculitic lesions, psoriasis, and lichenoid drug reactions.\textsuperscript{45}

**Protease inhibitors (ritonavir/lopinavir)**

These agents inhibit 3-chymotrypsin-like protease. Cutaneous adverse events of antiretroviral drugs include maculopapular drug eruptions, exfoliative erythroderma, Stevens-Johnson syndrome or toxic epidermal necrolysis.\textsuperscript{45}

**Ivermectin**

It has antimicrobial, anti-cancer properties and antiviral properties against wide range of viruses including SARS-CoV2. It acts by inhibiting the host importin alpha/beta-1 nuclear transport proteins, which are part of a key intracellular transport process that viruses hijack to enhance infection by suppressing the host antiviral response.\textsuperscript{45}

Some of the cutaneous side-effects include, swelling of ankles and hands and allergic skin reactions such as urticaria, rash, redness.

**Steroids**

Short term Corticosteroids have anti-inflammatory functions and they can suppress the inflammation during COVID-19 associated acute respiratory distress syndrome. Skin side-effects of steroids include folliculitis, acneiform eruptions, skin atrophy, telangiectasia, erythema, edema, acne, hirsutism, and stria.\textsuperscript{45}

**Barcitinib**

Janus kinase (JAK) inhibitors block viral entry into pneumocytes and can inhibit inflammatory mechanisms in COVID-19 infection. Palmoplantar pustulosis-like eruption, allergic skin rashes, herpes simplex, and herpes zoster infections, melanoma and nonmelanoma skin cancers, urticaria, rash, angioedema are some of the cutaneous side effects.\textsuperscript{45}

**Tocilizumab**

Tocilizumab, a humanized monoclonal antibody against IL-6 receptors, which is the chief cytokine responsible for the uncontrolled cytokine and chemokine response known as a “cytokine storm,” and this condition leads to the over-activation of effector T cells and production of pro-inflammatory cytokines. Skin infection, pruritus, skin hypersensitivity reactions, psoriasiform dermatitis are the cutaneous side-effects of tocilizumab.\textsuperscript{45}

**Anti-IL 1 (Anakinra)**

IL-1 blockade with anakinra helps in checking hyper inflammation during COVID-19 infection. Cutaneous side-effects of anakinra include rash, injection-site reactions, and skin infections such as wound infection, cellulitis.\textsuperscript{45}

**Anti TNF-α biologicals**

Severity of COVID-19 corresponds to the Higher TNF-α levels; hence adalimumab has been tried with success in some patients. Cutaneous side-effects of anti-TNFs include infusion and injection site reactions, psoriasis and psoriasiform-like lesions, lupus-like syndromes, cutaneous vasculitis, cutaneous infections, eczematous reactions, lichenoid eruptions, granulomatous reactions, cutaneous lymphoma, epithelial skin cancers or melanoma.\textsuperscript{45}

**INTRAVENOUS IMMUNOGLOBULIN (IVIG)**

High-dose intravenous immunoglobulin (IVIG) collected from recovered Coronavirus-19 patients may protect against COVID-19 and strengthen the immune system of new severe and treatment-resistant patients. Skin adverse events of IVIG treatments include anaphylactic reaction, facial vasculitic rash, urticaria, maculopapular rashes, petechiae, eczema, erythema multiforme, and alopecia.\textsuperscript{45}

**Zinc**

Elemental zinc (50 mg) supplementation is also advised in mildly symptomatic cases of COVID-19. Increased intracellular zinc concentrations efficiently impair replication in a number of RNA viruses. Zinc along with chloroquine has been shown to enhance cytotoxicity and induce apoptosis.\textsuperscript{46}
There are no cutaneous adverse events reported thus far but long-term zinc supplementation can cause copper deficiency, hence it is advised to keep a close eye on signs of copper deficiency like generalised skin color dilution and Menke’s kinky hair disease in future.47

**Vitamin C**

It is an antioxidant with anti-inflammatory properties, influences cellular immunity and vascular integrity.48 There is potential role of high doses of vitamin C in ameliorating inflammation and vascular injury in serious cases of COVID-19 causing sepsis and acute respiratory distress syndrome (ARDS). Hence it is now recommended in mild symptomatic cases practicing quarantine at home. No side effects are reported so far, because of water soluble nature of vitamin C.49

**CONCLUSION**

In the era of COVID-19 pandemic, where we see more and more publications stating various unforeseen presentations of COVID-19 infection, there is limited research when it comes to its cutaneous manifestations. COVID-19 being a viral illness, its cutaneous manifestations are similar to any viral illness and more.

Lately, many patients present with dermatological manifestation seldom preceding and in majority during COVID-19 infection. The order of appearance of these manifestations in combination with other symptoms of COVID-19 have major diagnostic and prognostic value.

Many cutaneous adverse drug reactions to drugs advocated in COVID-19 treatment are also being reported and with the fast-developing vaccine against COVID-19, getting to know the side effects of every drug is paramount, so that every clinician will get an insight into what to expect and when to expect in terms of dermatological manifestation of COVID-19 and its drugs. Through this article, we tried to present a brief review of some of the current cutaneous abnormalities observed in patients of COVID-19 perse and due to the drugs advocated for its treatment.

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**REFERENCES**


